Appln. No. 09/441,140
Amendment dated August 2, 2006
Reply to Office action of June 2, 2006

IN THE CLAIMS

Delete claims 155, 161, 167, 173-176, 183-186, 192-195 and 201-204.

Please amend claim 177 as follows:

177 (Amended). The therapeutic composition of claim
210 or 211, wherein said genetically-engineered monoclonal
antibody is a single-chain antibody.

Please insert new claims 210-213 as follows:

- 210 (New). A therapeutic composition, comprising: a pharmaceutical formulation comprising
- (1) a pharmaceutically acceptable carrier and
- (2) (a) a genetically-engineered antibody that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, or
- (b) a fragment of the genetically-engineered antibody of (a) that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid,

wherein said genetically-engineered antibody is obtained from a monoclonal antibody that

- (i) inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid and
- (ii) is obtainable using residues 1-28 of beta-amyloid as an immunogen or recognizes an epitope within residues 1-28 of beta-amyloid.
- 211 (New). The therapeutic composition of claim 210, wherein said genetically-engineered antibody of (2)(a) inhibits aggregation of human beta-amyloid or maintains the

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solubility of soluble human beta-amyloid, or said fragment of (2) (b) inhibits aggregation of human beta-amyloid or maintains the solubility of soluble human beta-amyloid, and said genetically-engineered antibody of (2) (a) is obtained from a monoclonal antibody that inhibits aggregation of human beta-amyloid or maintains the solubility of soluble human beta-amyloid and said monoclonal antibody is obtainable using residues 1-28 of human beta-amyloid as an immunogen or recognizes an epitope within residues 1-28 of human beta-amyloid.

- 212 (New). A therapeutic composition, comprising:

 a pharmaceutical formulation comprising
- (1) a pharmaceutically acceptable carrier and
- (2) (a) a human monoclonal antibody that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, or
- (b) a fragment of the human monoclonal antibody of (a) that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid,

wherein said human monoclonal antibody is obtainable using residues 1-28 of beta-amyloid as an immunogen.

213 (New). The therapeutic composition of claim
212, wherein said human monoclonal antibody of (2)(a) inhibits
aggregation of human beta-amyloid or maintains the solubility
of soluble human beta-amyloid, or said fragment of (2)(b)
inhibits aggregation of human beta-amyloid or maintains the
solubility of soluble human beta-amyloid, and wherein said

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human monoclonal antibody of (a) is obtainable using residues
1-28 of human beta-amyloid as an immunogen.

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